AMENDMENT TO THE SPECIFICATION

Please replace the paragraph beginning on page 8, line 25 with the following paragraph in amended format:

These tablets are tested for dissolution in standard apparatus type 1 of United States Pharmacopoeia. A 2% solution of sodium laurylsulfate in 0.01M potassium dihydrogenophosphate pH 6.8 buffer is used as dissolution medium. The amount of carbamazepine dissolved is recorded vs. time by using a Hewlett Packard HP8452A spectrophotometer. ~~The curve is given in figure 1.~~